

The impact of the assimilation of ozone from the Tropospheric Emission Spectrometer on surface ozone across North America

M. Parrington¹, D. B. A. Jones¹, K. W. Bowman², A. M. Thompson³, D. W. Tarasick⁴, J. Merrill⁵, S. J. Oltmans⁶, T. Leblanc², J. C. Witte⁷

¹University of Toronto; ²Jet Propulsion Laboratory; ³Pennsylvania State University; ⁴Environment Canada, ⁵University of Rhode Island, ⁶NOAA Climate Monitoring and Diagnostics Laboratory, ⁷Goddard Space Flight Center

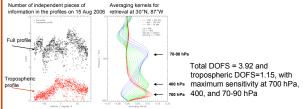
Abstract. We examine the potential of ozone data retrieved from the Tropospheric Emission Spectrometer (TES) for constraining background ozone values across North America in the GEOS-Chem chemical transport model. We assess here how changes in the abundance of background ozone in the free troposphere, following assimilation of TES data, impacts the surface ozone distribution in August 2006. We show that the assimilation cincreases the model surface ozone across the continent, with small changes of less than 3 ppbv in the eastern United States and larger changes of up to 9 ppbv over the western United States. This represents an increase of 15-25% of background ozone over the west coast and across Canada. We find that despite the good agreement between the assimilated ozone distribution and the ozonedonde measurements in the free tropsophere, comparisons with surface measurements from the EPA AQS and Environment Canada NAPS hewborks show that the assimilation exacerbates the basis insurface cozne in the model, suggesting potential model errors in the ozone precursor emissions or in the downward transport of ozone into the boundary layer in the ozone precursor emissions or or in the description of transport into the boundary layer in the boundary layer.

1. TES Instrument



- One of four instruments on the NASA EOS Aura spacecraft (launched July 15, 2004)
- Infrared Fourier transform spectrometer (3.3 - 15.4 um)
- Nadir footprint = 8 km x 5 km
- · Orbit repeats every 16 days
- Observations spaced about 2° along the orbit track
- Data products include O₃, CO, H₂O, HDO, and temperature

TES retrievals provide a smoothed representation of the true state



2. Chemical Data Assimilation Methodology

Sequential sub-optimal Kalman filter: $\hat{\mathbf{x}}^a = \mathbf{x}^f + \mathbf{K}[\mathbf{y}^{\text{obs}} - \mathbf{H}\mathbf{x}^f]$

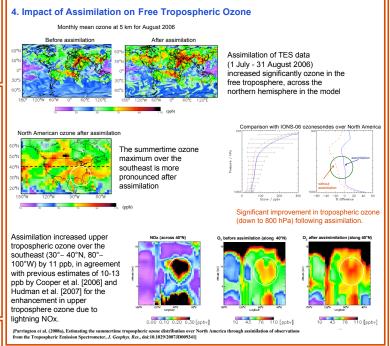
Kalman Gain Matrix: $\mathbf{K} = \mathbf{P}^f \mathbf{H}^T (\mathbf{H} \mathbf{P}^f \mathbf{H}^T + \mathbf{R})^{-1}$

Analysis Error Cov. Matrix: $\mathbf{P}^a = (\mathbf{I} - \mathbf{K}\mathbf{H})\mathbf{P}^f$

- Observation operator (H) accounts for TES averaging kernels and a priori profiles
- · Analysis error variance transported as a passive tracer
- O₃ and CO profile retrievals from TES
- 6-hour analysis cycle
- Assumed forecast error of 50% for CO and O₃
- Neglected horizontal correlations in forecast and observation error covariance matrices

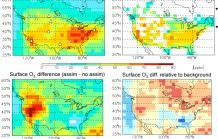
3. The GEOS-Chem Model

- · Chemical transport model
- 2.0° latitude x 2.5° longitude, 55 vertical levels (top level approx. 0.01 hPa)
- O₃-NOx-hydrocarbon chemistry
- Linearized (LINOZ) O₃ chemistry in the stratosphere
- Model transport driven by assimilated meteorological fields (GEOS-4) from NASA GMAO



5. Impact of Assimilation on Surface Ozone

Mean surface ozone (12h - 18h local time) for August 2006



2 3 4 5 6 7 8 9 [ppbv] 0 5 10 15 20 25 30

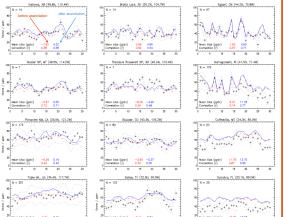
The model overestimates surface ozone in the east and underestimates it in the west Surface data from US Environmental Protection Agency Air Quality System (AQS) and the Environment Canada National Air Pollution Surveillance (NAPS) network, binned on the 2°x2.5° GEOSCHAPP (ACS)

 Increased free tropospheric ozone in the assimilation enhances the downward flux of background ozone into the North American boundary layer in the model

Surface ozone abundances in the assimilation increased by as much as 9 ppb, with the largest increase in western North America and the smallest increase in the southeastern USA.

6. Comparison with AQS and NAPS Ozone Data The model before assimilation is shown in red, whereas the model after assim

The model before assimilation is shown in red, whereas the model after assimilation is in blue. Surface data are shown indicated by the + symbols

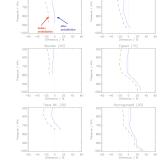


Assimilation reduced the bias in the western sites (which are influenced mainly by background ozone) and increased it in the east

Comparison with ozonesondes (August 2006)

The larger impact in the west is due, in part, to the higher elevation and deeper mixing layer in the west [e.g. Fiore et al. 2002; Dougherty 2008]

|Parrington et al. (2008b), The impact of the assimilation of ozone from the Tropospheric Emission Spectrometer on surface ozone across North America. Geophys. Res. Lett., submitted.]



7. Conclusions

- Assimilation of TES data reduced the bias in free tropospheric O₃ in the model from -35% to less than 5% (down to about 800 hPa)
- The improved distribution of O₃ in the free troposphere produced an increase in O₃ in the boundary layer of as much as 9 ppb (mainly in western North America)
- Increased background O₃ in the assimilation reduced the model bias in surface O₃ in western North America, but increased it in the east
- The increase in the bias in surface O_3 despite the improvement in free tropospheric O_3 indicates the presence of errors in the surface sources or sinks of O_3 or in the vertical transport of O_3 into the boundary layer in the model